DIAGNOSTIC REPORT

Patient Ref. No. 28000001068776

SRL Diagnostics

CLIENT CODE : C000138361

CLIENT'S NAME AND ADDRESS :
ACROFEMI HEALTHCARE LTD (MEDIWHEEL)
F-703, LADO SARAI, MEHRAULI
SOUTH WEST DELHI
NEW DELHI 110030
DELHI INDIA
8800465156

SRL Ltd
E-368, LGF, Nirman Vihar,Near Nirman Vihar Metro
NEW DELHI, 110092
NEW DELHI, INDIA
Tel : 9111591115, Fax :
CIN - U74899PB1995PLC045956
Email : wellness.eastdelhi@srl.in

PATIENT NAME : AJEET KUMAR SINGH PATIENT ID : **AJEEM10029028**

ACCESSION NO : **0028VE000539** AGE : 32 Years SEX : Male

DRAWN : RECEIVED : 14/05/2022 11:14 REPORTED : 17/05/2022 14:54

REFERRING DOCTOR : DR. MEDIWHEEL CLIENT PATIENT ID :

Test Report Status **Final**	Results		Biological Reference Interval	Units
MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE				
BLOOD COUNTS,EDTA WHOLE BLOOD				
HEMOGLOBIN	15.0		13.0 - 17.0	g/dL
METHOD : SPECTROPHOTOMETRY				
RED BLOOD CELL COUNT	5.28		4.5 - 5.5	mil/µL
METHOD : ELECTRICAL IMPEDANCE				
WHITE BLOOD CELL COUNT	8.00		4.0 - 10.0	thou/µL
METHOD : ELECTRICAL IMPEDANCE				
PLATELET COUNT	199		150 - 410	thou/µL
METHOD : ELECTRICAL IMPEDANCE				
RBC AND PLATELET INDICES				
HEMATOCRIT	45.6		40.0 - 50.0	%
METHOD : CALCULATED PARAMETER				
MEAN CORPUSCULAR VOL	86.3		83.0 - 101.0	fL
METHOD : DERIVED/COULTER PRINCIPLE				
MEAN CORPUSCULAR HGB.	28.4		27.0 - 32.0	pg
METHOD : CALCULATED PARAMETER				
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION	32.9		31.5 - 34.5	g/dL
METHOD : CALCULATED PARAMETER				
MENTZER INDEX	16.3			
RED CELL DISTRIBUTION WIDTH	**15.5**	**High**	11.6 - 14.0	%
METHOD : DERIVED/COULTER PRINCIPLE				
MEAN PLATELET VOLUME	9.8		6.8 - 10.9	fL
METHOD : DERIVED/COULTER PRINCIPLE				
WBC DIFFERENTIAL COUNT - NLR				
SEGMENTED NEUTROPHILS	66		40 - 80	%
METHOD : VCS TECHNOLOGY/ MICROSCOPY				
ABSOLUTE NEUTROPHIL COUNT	5.20		2.0 - 7.0	thou/µL
METHOD : CALCULATED PARAMETER				
LYMPHOCYTES	23		20 - 40	%
METHOD : VCS TECHNOLOGY/ MICROSCOPY				
ABSOLUTE LYMPHOCYTE COUNT	1.80		1.0 - 3.0	thou/µL
METHOD : CALCULATED PARAMETER				
NEUTROPHIL LYMPHOCYTE RATIO (NLR)	2.9			
EOSINOPHILS	2		1.0 - 6.0	%
METHOD : VCS TECHNOLOGY/ MICROSCOPY				

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ABSOLUTE EOSINOPHIL COUNT	0.16	0.02 - 0.50	thou/µL
METHOD : CALCULATED PARAMETER			
MONOCYTES	9	2.0 - 10.0	%
METHOD : VCS TECHNOLOGY/ MICROSCOPY			
ABSOLUTE MONOCYTE COUNT	0.70	0.2 - 1.0	thou/µL
METHOD : CALCULATED PARAMETER			
BASOPHILS	0	0 - 1	%
METHOD : VCS TECHNOLOGY/ MICROSCOPY			
DIFFERENTIAL COUNT PERFORMED ON:	EDTA SMEAR		
ERYTHRO SEDIMENTATION RATE, BLOOD			
SEDIMENTATION RATE (ESR)	14	0 - 14	mm at 1 hr
METHOD : WESTERGREN METHOD			
GLUCOSE, FASTING, PLASMA			
GLUCOSE, FASTING, PLASMA	94	74 - 106	mg/dL
METHOD : HEXOKINASE			
GLYCOSYLATED HEMOGLOBIN, EDTA WHOLE BLOOD			
GLYCOSYLATED HEMOGLOBIN (HBA1C)	5.4	Non-diabetic: < 5.7 Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5 ADA Target: 7.0 Action suggested: > 8.0	%
MEAN PLASMA GLUCOSE	108.3	< 116.0	mg/dL
GLUCOSE, POST-PRANDIAL, PLASMA			
GLUCOSE, POST-PRANDIAL, PLASMA	107	70 - 140	mg/dL
METHOD : HEXOKINASE			

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Test Report Status **Final**	Results	Biological Reference Interval	Units

Comments

Causes of Low Post Prandial Blood Sugar
The causes of low blood sugar that occurs following a meal have been divided traditionally into
(1)Alimentary
(2)Functional and
(3) That in diabetics and those with impaired glucose tolerance

Alimentary Hypoglycemia usually, but not necessarily, occurs in those patients who have had gastrointestinal surgery. Accelerated absorption of a glucose load leads to marked post - prandial hyperglycemia with a corresponding exaggerated insulin release thus resulting in hypoglycemia the ensuing hypoglycemia typically occurs from one and one half to three hours after eating. This pattern of glucose intolerance and a history of gastrointestinal surgery are suggestive of the diagnosis.

Functional Hypoglycemia is quite common in adults; it may be characterized by abnormally low plasma glucose and symptoms of light headedness, shakiness, diaphoresis, weakness, fatigue occurring with the modest withholding of food. Complaints suggestive of this syndrome commonly are seen in those who have emotional problems.

Post Prandial Hypoglycemia in diabetics: Patients who are diabetics or who have impaired glucose tolerance also may experience reactive hypoglycemia. In these patients, hypoglycemia occurs later than in the group with the alimentary disorder, and insulin response is delayed and exaggerated.

CORONARY RISK PROFILE (LIPID PROFILE), SERUM

CHOLESTEROL	134		< 200 Desirable 200 - 239 Borderline High >/= 240 High	mg/dL
METHOD : CHOLESTEROL OXIDASE, ESTERASE,PEROXIDASE				
TRIGLYCERIDES	92		< 150 Normal 150 - 199 Borderline High 200 - 499 High >/= 500 Very High	mg/dL
METHOD : ENZYMATIC, END POINT				
HDL CHOLESTEROL	**35**	**Low**	< 40 Low >/=60 High	mg/dL
METHOD : DIRECT MEASURE POLYMER-POLYANION				
DIRECT LDL CHOLESTEROL	79		< 100 Optimal 100 - 129 Near or above optimal 130 - 160 Borderline High 161 - 189 High >/= 190 Very High	mg/dL
METHOD : DIRECT MEASURE				
NON HDL CHOLESTEROL	99		Desirable: Less than 130 Above Desirable: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220	mg/dL
METHOD : CALCULATED PARAMETER				

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CHOL/HDL RATIO	3.8		3.3-4.4 Low Risk 4.5-7.0 Average Risk 7.1-11.0 Moderate Risk > 11.0 High Risk	
METHOD : CALCULATED PARAMETER				
LDL/HDL RATIO	2.3		0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk	
METHOD : CALCULATED PARAMETER				
VERY LOW DENSITY LIPOPROTEIN	18.4		Desirable value : 10 - 35	mg/dL
METHOD : CALCULATED PARAMETER				
LIVER FUNCTION PROFILE, SERUM				
BILIRUBIN, TOTAL	0.61		UPTO 1.2	mg/dL
METHOD : DIAZONIUM ION, BLANKED (ROCHE)				
BILIRUBIN, DIRECT	0.21		0.00 - 0.30	mg/dL
METHOD : DIAZOTIZATION				
BILIRUBIN, INDIRECT	0.40		0.00 - 0.60	mg/dL
METHOD : CALCULATED PARAMETER				
TOTAL PROTEIN	7.1		6.6 - 8.7	g/dL
METHOD : BIURET,SERUM BLANK,ENDPOINT				
ALBUMIN	4.5		3.97 - 4.94	g/dL
METHOD : BROMOCRESOL GREEN				
GLOBULIN	2.6		2.0 - 4.0 Neonates - Pre Mature: 0.29 - 1.04	g/dL
METHOD : CALCULATED PARAMETER				
ALBUMIN/GLOBULIN RATIO	1.7		1.0 - 2.0	RATIO
METHOD : CALCULATED PARAMETER				
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	34		0 - 40	U/L
METHOD : UV WITHOUT P5P				
ALANINE AMINOTRANSFERASE (ALT/SGPT)	**61**	**High**	0 - 41	U/L
METHOD : UV WITHOUT P5P				
ALKALINE PHOSPHATASE	102		40 - 129	U/L
METHOD : PNPP, AMP BUFFER-IFCC				
GAMMA GLUTAMYL TRANSFERASE (GGT)	33		8 - 61	U/L
METHOD : G-GLUTAMYL-CARBOXY-NITROANILIDE-IFCC				
LACTATE DEHYDROGENASE	197		135 - 225	U/L
METHOD : L TO P, IFCC				
SERUM BLOOD UREA NITROGEN				

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BLOOD UREA NITROGEN METHOD : UREASE - UV	15		6 - 20	mg/dL
CREATININE, SERUM				
CREATININE METHOD : ALKALINE PICRATE-KINETIC	0.85		0.70 - 1.20	mg/dL
BUN/CREAT RATIO				
BUN/CREAT RATIO METHOD : CALCULATED PARAMETER	**17.65**	**High**	5.00 - 15.00	
URIC ACID, SERUM				
URIC ACID METHOD : URICASE, COLORIMETRIC	6.1		3.4 - 7.0	mg/dL
TOTAL PROTEIN, SERUM				
TOTAL PROTEIN METHOD : BIURET,SERUM BLANK,ENDPOINT	7.1		6.6 - 8.7	g/dL
ALBUMIN, SERUM				
ALBUMIN METHOD : BROMOCRESOL GREEN	4.5		3.97 - 4.94	g/dL
GLOBULIN				
GLOBULIN METHOD : CALCULATED PARAMETER	2.6		2.0 - 4.0 Neonates - Pre Mature: 0.29 - 1.04	g/dL
ELECTROLYTES (NA/K/CL), SERUM				
SODIUM METHOD : ISE INDIRECT	137		136 - 145	mmol/L
POTASSIUM METHOD : ISE INDIRECT	3.81		3.5 - 5.1	mmol/L
CHLORIDE METHOD : ISE INDIRECT	101		98 - 107	mmol/L
PHYSICAL EXAMINATION, URINE				
COLOR METHOD : VISUAL	PALE YELLOW			
APPEARANCE METHOD : VISUAL	CLEAR			
SPECIFIC GRAVITY METHOD : PKA CHANGE OF PRETREATED POLYELECTROLYTES	<=1.005		1.003 - 1.035	
CHEMICAL EXAMINATION, URINE				

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PH	6.0	4.7 - 7.5	
METHOD : DOUBLE INDICATOR PRINCIPLE			
PROTEIN	NOT DETECTED	NOT DETECTED	
METHOD : PROTEIN- ERROR INDICATOR			
GLUCOSE	NOT DETECTED		
METHOD : OXIDASE-PEROXIDASE REACTION			
KETONES	NOT DETECTED	NOT DETECTED	
METHOD : ACETOACETIC REACTION WITH NITROPRUSSIDE			
BLOOD	NOT DETECTED	NOT DETECTED	
METHOD : PEROXIDASE-LIKE ACTIVITY OF HEMOGLOBIN			
BILIRUBIN	NOT DETECTED	NOT DETECTED	
METHOD : DIAZOTIZATION			
UROBILINOGEN	NORMAL	NORMAL	
METHOD : MODIFIED EHRLICH REACTION			
NITRITE	NOT DETECTED	NOT DETECTED	
METHOD : CONVERTION OF NITRATE TO NITRITE			
MICROSCOPIC EXAMINATION, URINE			
PUS CELL (WBC'S)	3-5	0-5	/HPF
METHOD : MICROSCOPIC EXAMINATION			
EPITHELIAL CELLS	0-1	0-5	/HPF
METHOD : MICROSCOPIC EXAMINATION			
ERYTHROCYTES (RBC'S)	NOT DETECTED	NOT DETECTED	/HPF
METHOD : MICROSCOPIC EXAMINATION			
CASTS	NOT DETECTED		
METHOD : MICROSCOPIC EXAMINATION			
CRYSTALS	NOT DETECTED		
METHOD : MICROSCOPIC EXAMINATION			
BACTERIA	**DETECTED (FEW)**	NOT DETECTED	
METHOD : MICROSCOPIC EXAMINATION			
YEAST	NOT DETECTED	NOT DETECTED	
REMARKS	MICROSCOPIC EXAMINATION DONE ON CENTRIFUGED URINE PLEASE NOTE THAT GRADING OF BACTERIA NEEDS TO BE CORELATED WITH THE CULTURE IN CASE FOUND SIGNIFICANT CLINICALLY. THE BACTERIA CAN ALSO BE A PART OF SURROUNDING SKIN FLORA.		
METHOD : MANUAL			
THYROID PANEL, SERUM			
T3	137.0	80.00 - 200.00	ng/dL
METHOD : ECLIA			

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T4	8.66	5.10 - 14.10	µg/dL
METHOD : ECLIA			
TSH 3RD GENERATION	2.660	0.270 - 4.200	µIU/mL
METHOD : ECLIA			

Comments

Note: TSH
Please note that TSH values can show a diurnal variation (up to 50%). Subclinical thyroid/ gut diseases/ intake of calcium, multivitamin and several other drugs, resistance to thyroid hormones, noncompliance to medication can lead to discrepant thyroid results, the levels of thyroid can also be affected by weather, high fiber diet, estrogen surge, stress, alcohol, pregnancy, obesity/weight loss. The results can vary on different instruments. Serum TSH changes significantly in response to even minor changes in thyroid hormones. For the diagnosis of hypo and hyper thyroids, sole dependence on TSH should not be done and testing also needs to be performed for T3, T4 and other metabolic parameters as well as reasons elicited above.

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD

ABO GROUP	TYPE A
METHOD : COLUMN AGGLUTINATION TECHNOLOGY/AGGLUTINATION TECHNOLOGY	
RH TYPE	POSITIVE
METHOD : COLUMN AGGLUTINATION TECHNOLOGY/AGGLUTINATION TECHNOLOGY	

XRAY-CHEST

»»	BOTH THE LUNG FIELDS ARE CLEAR
»»	BOTH THE COSTOPHRENIC AND CARIOPHRENIC ANGELS ARE CLEAR
»»	BOTH THE HILA ARE NORMAL
»»	CARDIAC AND AORTIC SHADOWS APPEAR NORMAL
»»	BOTH THE DOMES OF THE DIAPHRAM ARE NORMAL
»»	VISUALIZED BONY THORAX IS NORMAL
IMPRESSION	NO ABNORMALITY DETECTED

TMT OR ECHO

TMT OR ECHO	PENDING

ECG

ECG	WITHIN NORMAL LIMITS

MEDICAL HISTORY

RELEVANT PRESENT HISTORY	NOT SIGNIFICANT
RELEVANT PAST HISTORY	NOT SIGNIFICANT
RELEVANT PERSONAL HISTORY	MARRIED,NON VEGETARIAN
RELEVANT FAMILY HISTORY	FATHER-DIABETS MELLITUS
OCCUPATIONAL HISTORY	NOT SIGNIFICANT
HISTORY OF MEDICATIONS	NOT SIGNIFICANT

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ANTHROPOMETRIC DATA & BMI			
HEIGHT IN METERS	1.72		mts
WEIGHT IN KGS.	92.8		Kgs
BMI	31	BMI & Weight Status as follows: Below 18.5: Underweight 18.5 - 24.9: Normal 25.0 - 29.9: Overweight 30.0 and Above: Obese	kg/sqmts
GENERAL EXAMINATION			
MENTAL / EMOTIONAL STATE	NORMAL		
PHYSICAL ATTITUDE	NORMAL		
GENERAL APPEARANCE / NUTRITIONAL STATUS	OVERWEIGHT		
BUILT / SKELETAL FRAMEWORK	AVERAGE		
FACIAL APPEARANCE	NORMAL		
SKIN	NORMAL		
UPPER LIMB	NORMAL		
LOWER LIMB	NORMAL		
NECK	NORMAL		
NECK LYMPHATICS / SALIVARY GLANDS	NOT ENLARGED OR TENDER		
THYROID GLAND	NOT ENLARGED		
CAROTID PULSATION	NORMAL		
TEMPERATURE	NORMAL		
PULSE	62 / MIN REGULAR, ALL PERIPHERAL PULSES WELL FELT, NO CAROTID BRUIT		
RESPIRATORY RATE	NORMAL		
CARDIOVASCULAR SYSTEM			
BP	110/80		mm/Hg
PERICARDIUM	NORMAL		
APEX BEAT	NORMAL		
HEART SOUNDS	S1, S2 HEARD NORMALLY		
MURMURS	ABSENT		
RESPIRATORY SYSTEM			
SIZE AND SHAPE OF CHEST	NORMAL		
MOVEMENTS OF CHEST	SYMMETRICAL		
BREATH SOUNDS INTENSITY	NORMAL		
BREATH SOUNDS QUALITY	VESICULAR (NORMAL)		

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ADDED SOUNDS	ABSENT		
PER ABDOMEN			
APPEARANCE	NORMAL		
VENOUS PROMINENCE	ABSENT		
LIVER	NOT PALPABLE		
SPLEEN	NOT PALPABLE		
CENTRAL NERVOUS SYSTEM			
HIGHER FUNCTIONS	NORMAL		
CRANIAL NERVES	NORMAL		
CEREBELLAR FUNCTIONS	NORMAL		
SENSORY SYSTEM	NORMAL		
MOTOR SYSTEM	NORMAL		
REFLEXES	NORMAL		
MUSCULOSKELETAL SYSTEM			
SPINE	NORMAL		
JOINTS	NORMAL		
BASIC EYE EXAMINATION			
CONJUNCTIVA	NORMAL		
EYELIDS	NORMAL		
EYE MOVEMENTS	NORMAL		
CORNEA	NORMAL		
DISTANT VISION RIGHT EYE WITHOUT GLASSES	NORMAL		
DISTANT VISION LEFT EYE WITHOUT GLASSES	NORMAL		
NEAR VISION RIGHT EYE WITHOUT GLASSES	NORMAL		
NEAR VISION LEFT EYE WITHOUT GLASSES	NORMAL		
COLOUR VISION	NORMAL		
BASIC ENT EXAMINATION			
EXTERNAL EAR CANAL	NORMAL		
TYMPANIC MEMBRANE	NORMAL		
NOSE	NO ABNORMALITY DETECTED		
SINUSES	CLEAR		
THROAT	NO ABNORMALITY DETECTED		
TONSILS	NOT ENLARGED		
SUMMARY			

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DRAWN : RECEIVED : 14/05/2022 11:14 REPORTED : 17/05/2022 14:54

REFERRING DOCTOR : DR. MEDIWHEEL CLIENT PATIENT ID :

Test Report Status **Final**	Results	Biological Reference Interval	Units
RELEVANT HISTORY	NOT SIGNIFICANT		
RELEVANT GP EXAMINATION FINDINGS	NOT SIGNIFICANT		
RELEVANT NON PATHOLOGY DIAGNOSTICS	CHEST X RAY PA-NO ABNORMALITIES DETECTED		
REMARKS / RECOMMENDATIONS	FOLLOW UP TO MD PHYSICIAN WITH ALL REPORTS FOR FURTHER MANAGEMENT. GENERAL PHYSICAL EXAMINATION IS NORMAL."		

Interpretation(s)
BLOOD COUNTS,EDTA WHOLE BLOOD-
The cell morphology is well preserved for 24hrs. However after 24-48 hrs a progressive increase in MCV and HCT is observed leading to a decrease in MCHC. A direct smear is recommended for an accurate differential count and for examination of RBC morphology.
RBC AND PLATELET INDICES-
Mentzer index (MCV/RBC) is an automated cell-counter based calculated screen tool to differentiate cases of Iron deficiency anaemia(>13) from Beta thalassaemia trait (<13) in patients with microcytic anaemia. This needs to be interpreted in line with clinical correlation and suspicion. Estimation of HbA2 remains the gold standard for diagnosing a case of beta thalassaemia trait.
WBC DIFFERENTIAL COUNT - NLR-
The optimal threshold of 3.3 for NLR showed a prognostic possibility of clinical symptoms to change from mild to severe in COVID positive patients. When age = 49.5 years old and NLR = 3.3, 46.1% COVID-19 patients with mild disease might become severe. By contrast, when age < 49.5 years old and NLR < 3.3, COVID-19 patients tend to show mild disease.
(Reference to - The diagnostic and predictive role of NLR, d-NLR and PLR in COVID-19 patients ; A.-P. Yang, et al.; International Immunopharmacology 84 (2020) 106504
This ratio element is a calculated parameter and out of NABL scope.
ERYTHRO SEDIMENTATION RATE, BLOOD-
Erythrocyte sedimentation rate (ESR) is a non - specific phenomena and is clinically useful in the diagnosis and monitoring of disorders associated with an increased production of acute phase reactants. The ESR is increased in pregnancy from about the 3rd month and returns to normal by the 4th week post partum. ESR is influenced by age, sex, menstrual cycle and drugs (eg. corticosteroids, contraceptives). It is especially low (0 -1mm) in polycythaemia, hypofibrinogenemia or congestive cardiac failure and when there are abnormalities of the red cells such as poikilocytosis, spherocytosis or sickle cells.

Reference :
1. Nathan and Oski's Haematology of Infancy and Childhood, 5th edition
2. Paediatric reference intervals. AACC Press, 7th edition. Edited by S. Soldin
3. The reference for the adult reference range is "Practical Haematology by Dacie and Lewis, 10th Edition"
GLUCOSE, FASTING, PLASMA-
ADA 2021 guidelines for adults, after 8 hrs fasting is as follows:
Pre-diabetics: 100 - 125 mg/dL
Diabetic: > or = 126 mg/dL
GLYCOSYLATED HEMOGLOBIN, EDTA WHOLE BLOOD-
Glycosylated hemoglobin (GHb) has been firmly established as an index of long-term blood glucose concentrations and as a measure of the risk for the development of complications in patients with diabetes mellitus. Formation of GHb is essentially irreversible, and the concentration in the blood depends on both the life span of the red blood cell (average 120 days) and the blood glucose concentration. Because the rate of formation of GHb is directly proportional to the concentration of glucose in the blood, the GHb concentration represents the integrated values for glucose over the preceding 6-8 weeks.
Any condition that alters the life span of the red blood cells has the potential to alter the GHb level. Samples from patients with hemolytic anemias will exhibit decreased glycated hemoglobin values due to the shortened life span of the red cells. This effect will depend upon the severity of the anemia. Samples from patients with polycythemia or post-splenectomy may exhibit increased glycated hemoglobin values due to a somewhat longer life span of the red cells.
Glycosylated hemoglobins results from patients with HbSS, HbCC, and HbSC and HbD must be interpreted with caution, given the pathological processes, including anemia, increased red cell turnover, transfusion requirements, that adversely impact HbA1c as a marker of long-term glycemic control. In these conditions, alternative forms of testing such as glycated serum protein (fructosamine) should be considered.
"Targets should be individualized; More or less stringent glycemic goals may be appropriate for individual patients. Goals should be individualized based on duration of diabetes, age/life expectancy, comorbid conditions, known CVD or advanced microvascular complications, hypoglycemia unawareness, and individual patient considerations."

References
1. Tietz Textbook of Clinical Chemistry and Molecular Diagnostics, edited by Carl A Burtis, Edward R.Ashwood, David E Bruns, 4th Edition, Elsevier publication, 2006, 879-884.
2. Forsham PH. Diabetes Mellitus:A rational plan for management. Postgrad Med 1982, 71,139-154.
3. Mayer TK, Freedman ZR: Protein glycosylation in Diabetes Mellitus: A review of laboratory measurements and their clinical utility. Clin Chim Acta 1983, 127, 147-184.
GLUCOSE, POST-PRANDIAL, PLASMA-ADA Guidelines for 2hr post prandial glucose levels is only after ingestion of 75grams of glucose in 300 ml water,over a period of 5 minutes.
CORONARY RISK PROFILE (LIPID PROFILE), SERUM.-

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DIAGNOSTIC REPORT

Patient Ref. No. 28000001068776

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Serum cholesterol is a blood test that can provide valuable information for the risk of coronary artery disease This test can help determine your risk of the build up of plaques in your arteries that can lead to narrowed or blocked arteries throughout your body (atherosclerosis). High cholesterol levels usually don't cause any signs or symptoms, so a cholesterol test is an important tool. High cholesterol levels often are a significant risk factor for heart disease and important for diagnosis of hyperlipoproteinemia, atherosclerosis, hepatic and thyroid diseases.

Serum Triglyceride are a type of fat in the blood. When you eat, your body converts any calories it doesn't need into triglycerides, which are stored in fat cells. High triglyceride levels are associated with several factors, including being overweight, eating too many sweets or drinking too much alcohol, smoking, being sedentary, or having diabetes with elevated blood sugar levels. Analysis has proven useful in the diagnosis and treatment of patients with diabetes mellitus, nephrosis, liver obstruction, other diseases involving lipid metabolism, and various endocrine disorders. In conjunction with high density lipoprotein and total serum cholesterol, a triglyceride determination provides valuable information for the assessment of coronary heart disease risk.It is done in fasting state.

High-density lipoprotein (HDL) cholesterol. This is sometimes called the ""good"" cholesterol because it helps carry away LDL cholesterol, thus keeping arteries open and blood flowing more freely.HDL cholesterol is inversely related to the risk for cardiovascular disease. It increases following regular exercise, moderate alcohol consumption and with oral estrogen therapy. Decreased levels are associated with obesity, stress, cigarette smoking and diabetes mellitus.

SERUM LDL The small dense LDL test can be used to determine cardiovascular risk in individuals with metabolic syndrome or established/progressing coronary artery disease, individuals with triglyceride levels between 70 and 140 mg/dL, as well as individuals with a diet high in trans-fat or carbohydrates. Elevated sdLDL levels are associated with metabolic syndrome and an 'atherogenic lipoprotein profile', and are a strong, independent predictor of cardiovascular disease.
Elevated levels of LDL arise from multiple sources. A major factor is sedentary lifestyle with a diet high in saturated fat. Insulin-resistance and pre-diabetes have also been implicated, as has genetic predisposition. Measurement of sdLDL allows the clinician to get a more comprehensive picture of lipid risk factors and tailor treatment accordingly. Reducing LDL levels will reduce the risk of CVD and MI.

Non HDL Cholesterol - Adult treatment panel ATP III suggested the addition of Non-HDL Cholesterol as an indicator of all atherogenic lipoproteins (mainly LDL and VLDL). NICE guidelines recommend Non-HDL Cholesterol measurement before initiating lipid lowering therapy. It has also been shown to be a better marker of risk in both primary and secondary prevention studies.

Recommendations:
Results of Lipids should always be interpreted in conjunction with the patient's medical history, clinical presentation and other findings.

NON FASTING LIPID PROFILE includes Total Cholesterol, HDL Cholesterol and calculated non-HDL Cholesterol. It does not include triglycerides and may be best used in patients for whom fasting is difficult.
LIVER FUNCTION PROFILE, SERUM-
LIVER FUNCTION PROFILE
Bilirubin is a yellowish pigment found in bile and is a breakdown product of normal heme catabolism. Bilirubin is excreted in bile and urine, and elevated levels may give yellow discoloration in jaundice.Elevated levels results from increased bilirubin production (eg, hemolysis and ineffective erythropoiesis), decreased bilirubin excretion (eg, obstruction and hepatitis), and abnormal bilirubin metabolism (eg, hereditary and neonatal jaundice). Conjugated (direct) bilirubin is elevated more than unconjugated (indirect) bilirubin in Viral hepatitis, Drug reactions, Alcoholic liver disease Conjugated (direct) bilirubin is also elevated more than unconjugated (indirect) bilirubin when there is some kind of blockage of the bile ducts like in Gallstones getting into the bile ducts, tumors &Scarring of the bile ducts. Increased unconjugated (indirect) bilirubin may be a result of Hemolytic or pernicious anemia, Transfusion reaction & a common metabolic condition termed Gilbert syndrome, due to low levels of the enzyme that attaches sugar molecules to bilirubin.
AST is an enzyme found in various parts of the body. AST is found in the liver, heart, skeletal muscle, kidneys, brain, and red blood cells, and it is commonly measured clinically as a marker for liver health. AST levels increase during chronic viral hepatitis, blockage of the bile duct, cirrhosis of the liver,liver cancer,kidney failure,hemolytic anemia,pancreatitis,hemochromatosis. AST levels may also increase after a heart attack or strenuous activity.ALT test measures the amount of this enzyme in the blood.ALT is found mainly in the liver, but also in smaller amounts in the kidneys,heart,muscles, and pancreas.It is commonly measured as a part of a diagnostic evaluation of hepatocellular injury, to determine liver health.AST levels increase during acute hepatitis,sometimes due to a viral infection,ischemia to the liver,chronic hepatitis,obstruction of bile ducts,cirrhosis.
ALP is a protein found in almost all body tissues.Tissues with higher amounts of ALP include the liver,bile ducts and bone.Elevated ALP levels are seen in Biliary obstruction, Osteoblastic bone tumors, osteomalacia, hepatitis, Hyperparathyroidism, Leukemia, Lymphoma, Paget's disease,Rickets,Sarcoidosis etc. Lower-than-normal ALP levels seen in Hypophosphatasia,Malnutrition,Protein deficiency,Wilson's disease.GGT is an enzyme found in cell membranes of many tissues mainly in the liver,kidney and pancreas.It is also found in other tissues including intestine,spleen,heart, brain and seminal vesicles.The highest concentration is in the kidney,but the liver is considered the source of normal enzyme activity.Serum GGT has been widely used as an index of liver dysfunction.Elevated serum GGT activity can be found in diseases of the liver,biliary system and pancreas.Conditions that increase serum GGT are obstructive liver disease,high alcohol consumption and use of enzyme-inducing drugs etc.Serum total protein,also known as total protein,is a biochemical test for measuring the total amount of protein in serum.Protein in the plasma is made up of albumin and globulin.Higher-than-normal levels may be due to:Chronic inflammation or infection,including HIV and hepatitis B or C,Multiple myeloma,Waldenstrom's disease.Lower-than-normal levels may be due to: Agammaglobulinemia,Bleeding (hemorrhage),Burns,Glomerulonephritis,Liver disease, Malabsorption,Malnutrition,Nephrotic syndrome,Protein-losing enteropathy etc.Human serum albumin is the most abundant protein in human blood plasma.It is produced in the liver.Albumin constitutes about half of the blood serum protein.Low blood albumin levels (hypoalbuminemia) can be caused by:Liver disease like cirrhosis of the liver, nephrotic syndrome,protein-losing enteropathy,Burns,hemodilution,increased vascular permeability or decreased lymphatic clearance,malnutrition and wasting etc
SERUM BLOOD UREA NITROGEN-
Causes of Increased levels
Pre renal
• High protein diet, Increased protein catabolism, GI haemorrhage, Cortisol, Dehydration, CHF Renal
• Renal Failure
Post Renal
• Malignancy, Nephrolithiasis, Prostatism

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Causes of decreased levels
• Liver disease
• SIADH.
CREATININE, SERUM-
Higher than normal level may be due to:
• Blockage in the urinary tract
• Kidney problems, such as kidney damage or failure, infection, or reduced blood flow
• Loss of body fluid (dehydration)
• Muscle problems, such as breakdown of muscle fibers
• Problems during pregnancy, such as seizures (eclampsia)), or high blood pressure caused by pregnancy (preeclampsia)

Lower than normal level may be due to:
• Myasthenia Gravis
• Muscular dystrophy
URIC ACID, SERUM-
Causes of Increased levels
Dietary
• High Protein Intake.
• Prolonged Fasting,
• Rapid weight loss.
Gout
Lesch nyhan syndrome.
Type 2 DM.
Metabolic syndrome.

Causes of decreased levels
• Low Zinc Intake
• OCP's
• Multiple Sclerosis

Nutritional tips to manage increased Uric acid levels
• Drink plenty of fluids
• Limit animal proteins
• High Fibre foods
• Vit C Intake
• Antioxidant rich foods
TOTAL PROTEIN, SERUM-
Serum total protein,also known as total protein, is a biochemical test for measuring the total amount of protein in serum..Protein in the plasma is made up of albumin and globulin

Higher-than-normal levels may be due to: Chronic inflammation or infection, including HIV and hepatitis B or C, Multiple myeloma, Waldenstrom's disease
Lower-than-normal levels may be due to: Agammaglobulinemia, Bleeding (hemorrhage),Burns,Glomerulonephritis, Liver disease, Malabsorption, Malnutrition, Nephrotic syndrome,Protein-losing enteropathy etc.
ALBUMIN, SERUM-
Human serum albumin is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. Low blood albumin levels (hypoalbuminemia) can be caused by: Liver disease like cirrhosis of the liver, nephrotic syndrome, protein-losing enteropathy, Burns, hemodilution, increased vascular permeability or decreased lymphatic clearance,malnutrition and wasting etc.
ELECTROLYTES (NA/K/CL), SERUM-
Sodium levels are Increased in dehydration, cushing's syndrome, aldosteronism & decreased in Addison's disease, hypopituitarism,liver disease. Hypokalemia (low K) is common in vomiting, diarrhea, alcoholism, folic acid deficiency and primary aldosteronism. Hyperkalemia may be seen in end-stage renal failure, hemolysis, trauma, Addison's disease, metabolic acidosis, acute starvation, dehydration, and with rapid K infusion.Chloride is increased in dehydration, renal tubular acidosis (hyperchloremia metabolic acidosis), acute renal failure, metabolic acidosis associated with prolonged diarrhea and loss of sodium bicarbonate, diabetes insipidus, adrenocortical hyperfuction, salicylate intoxication and with excessive infusion of isotonic saline or extremely high dietary intake of salt.Chloride is decreased in overhydration, chronic respiratory acidosis, salt-losing nephritis, metabolic alkalosis, congestive heart failure, Addisonian crisis, certain types of metabolic acidosis, persistent gastric secretion and prolonged vomiting,
THYROID PANEL, SERUM-
Triiodothyronine T3 , is a thyroid hormone. It affects almost every physiological process in the body, including growth, development, metabolism, body temperature, and heart rate. Production of T3 and its prohormone thyroxine (T4) is activated by thyroid-stimulating hormone (TSH), which is released from the pituitary gland. Elevated concentrations of T3, and T4 in the blood inhibit the production of TSH.
Thyroxine T4, Thyroxine's principal function is to stimulate the metabolism of all cells and tissues in the body. Excessive secretion of thyroxine in the body is hyperthyroidism, and deficient secretion is called hypothyroidism. Most of the thyroid hormone in blood is bound to transport proteins. Only a very small fraction of the circulating hormone is free and biologically active.
In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiary hypothyroidism, TSH levels are low.
Below mentioned are the guidelines for Pregnancy related reference ranges for Total T4, TSH & Total T3

Levels in Pregnancy	TOTAL T4 (µg/dL)	TSH3G (µIU/mL)	TOTAL T3 (ng/dL)

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First Trimester	6.6 - 12.4	0.1 - 2.5	81 - 190
2nd Trimester	6.6 - 15.5	0.2 - 3.0	100 - 260
3rd Trimester	6.6 - 15.5	0.3 - 3.0	100 - 260

Below mentioned are the guidelines for age related reference ranges for T3 and T4.

	T3 (ng/dL)	T4 (µg/dL)
New Born:	75 - 260	1-3 day: 8.2 - 19.9
.		1 Week: 6.0 - 15.9

NOTE: TSH concentrations in apparently normal euthyroid subjects are known to be highly skewed, with a strong tailed distribution towards higher TSH values. This is well documented in the pediatric population including the infant age group.
Kindly note: Method specific reference ranges are appearing on the report under biological reference range.

Reference:
1. Burtis C.A., Ashwood E. R. Bruns D.E. Teitz textbook of Clinical Chemistry and Molecular Diagnostics, 4th Edition.
2. Gowenlock A.H. Varley's Practical Clinical Biochemistry, 6th Edition.
3. Behrman R.E. Kilegman R.M., Jenson H. B. Nelson Text Book of Pediatrics, 17th Edition
ABO GROUP & RH TYPE, EDTA WHOLE BLOOD-
Blood group is identified by antigens and antibodies present in the blood. Antigens are protein molecules found on the surface of red blood cells. Antibodies are found in plasma. To determine blood group, red cells are mixed with different antibody solutions to give A,B,O or AB.

Disclaimer: "Please note, as the results of previous ABO and Rh group (Blood Group) for pregnant women are not available, please check with the patient records for availability of the same."

The test is performed by both forward as well as reverse grouping methods.
MEDICAL
HISTORY-**********
THIS REPORT CARRIES THE SIGNATURE OF OUR LABORATORY DIRECTOR. THIS IS AN INVIOLABLE FEATURE OF OUR LAB MANAGEMENT SOFTWARE. HOWEVER, ALL EXAMINATIONS AND INVESTIGATIONS HAVE BEEN CONDUCTED BY OUR PANEL OF DOCTORS.

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MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE

ULTRASOUND ABDOMEN

ULTRASOUND ABDOMEN

PENDING

****End Of Report****
Please visit www.srlworld.com for related Test Information for this accession

Dr. Shyla Goel,M.B.B.S ,DCP .
Pathologist .

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2. All Tests are performed and reported as per the turnaround time stated in the SRL Directory of services (DOS).
3. SRL confirms that all tests have been performed or assayed with highest quality standards, clinical safety & technical integrity.
4. A requested test might not be performed if:
   a. Specimen received is insufficient or inappropriate specimen quality is unsatisfactory
   b. Incorrect specimen type
   c. Request for testing is withdrawn by the ordering doctor or patient
   d. There is a discrepancy between the label on the specimen container and the name on the test requisition form
5. The results of a laboratory test are dependent on the quality of the sample as well as the assay technology.
6. Result delays could be because of uncontrolled circumstances. e.g. assay run failure.
7. Tests parameters marked by asterisks are excluded from the "scope" of NABL accredited tests. (If laboratory is accredited).
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